

## STUDYING OZONOLYSIS REACTIONS OF 2-BUTENES USING CAVITY RING-DOWN SPECTROSCOPY

LIMING WANG, *School of Chemistry and Chemical Engineering, South China University of Technology, Guangzhou, China*; YINGDI LIU, *Jet Propulsion Laboratory, California Institute of Technology, Pasadena, CA, USA*; MIXTLI CAMPOS-PINEDA, CHAD PRIEST, *Department of Chemistry, University of California, Riverside, CA, USA*; JINGSONG ZHANG, *Department of Chemistry and Air Pollution Research Center, University of California, Riverside, CA, USA*.

Ozonolysis reactions of alkenes are important oxidation pathways of alkenes in the atmosphere, and they are also significant sources of tropospheric hydroxyl radicals. In this work, ozone reactions with *trans*- and *cis*-2-butene are studied using cavity ring-down spectroscopy (CRDS). Vinoxyl ( $\text{CH}_2\text{CHO}$ ) radical, a proposed co-product of OH from dissociation of Criegee intermediates following the primary ozonolysis of 2-butenes, is directly observed. The vinoxy formation is found to decrease with increasing pressure. Formaldehyde ( $\text{HCHO}$ ), a side-product in the ozonolysis reactions, is also monitored. Chemical kinetic modeling has been performed to illustrate the reaction mechanisms and to quantify the reactive intermediate and product yields.